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| NEWS | 4 | JAN | 16 | IPC version 2007.01 thesaurus available on STN |
| NEWS | 5 | JAN | 16 | WPIDS/WPINDEX/WPIX enhanced with IPC 8 reclassification data |
| NEWS | 6 | JAN | 22 | CA/CAplus updated with revised CAS roles |
| NEWS | 7 | JAN | 22 | CA/CAplus enhanced with patent applications from India |
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| NEWS | 17 | FEB | 26 | CAS Registry Number crossover limit increased from 10,000 to 300,000 in multiple databases |
| NEWS | 18 | MAR | 15 | WPIDS/WPIX enhanced with new FRAGHITSTR display format |
| NEWS | 19 | MAR | 16 | CASREACT coverage extended |
| NEWS | 20 | MAR | 20 | MARPAT now updated daily |
| NEWS | 21 | MAR | 22 | LWPI reloaded |
| NEWS | 22 | MAR | 30 | RDISCLOSURE reloaded with enhancements |
| NEWS | 23 | APR | 02 | JICST-EPLUS removed from database clusters and STN |
| NEWS | 24 | APR | 30 | GENBANK reloaded and enhanced with Genome Project ID field |
| NEWS | 25 | APR | 30 | CHEMCATS enhanced with 1.2 million new records |
| NEWS | 26 | APR | 30 | CA/CAplus enhanced with 1870-1889 U.S. patent records |
| NEWS | 27 | APR | 30 | INPADOC replaced by INPADOCDB on STN |
| NEWS | 28 | MAY | 01 | New CAS web site launched |
| NEWS | 29 | MAY | 08 | CA/CAplus Indian patent publication number format defined |
| NEWS | 30 | MAY | 14 | RDISCLOSURE on STN Easy enhanced with new search and display fields |
| NEWS | 31 | MAY | 21 | BIOSIS reloaded and enhanced with archival data |
| NEWS | 32 | MAY | 21 | TOXCENTER enhanced with BIOSIS reload |
| NEWS | 33 | MAY | 21 | CA/CAplus enhanced with additional kind codes for German patents |
| NEWS | 34 | MAY | 22 | CA/CAplus enhanced with IPC reclassification in Japanese patents |
| NEWS EXPRESS | | NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006. | | |
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| NEWS IPC8 | | For general information regarding STN implementation of IPC 8 | | |

Enter NEWS followed by the item number or name to see news on that specific topic.

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FILE 'HOME' ENTERED AT 15:51:36 ON 15 JUN 2007

FILE 'MEDLINE' ENTERED AT 15:51:56 ON 15 JUN 2007

FILE 'BIOSIS' ENTERED AT 15:51:56 ON 15 JUN 2007
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=> (triacylglycerol production)
(TRIACYLGLYCEROL IS NOT A RECOGNIZED COMMAND
The previous command name entered was not recognized by the system.
For a list of commands available to you in the current file, enter
"HELP COMMANDS" at an arrow prompt (>).

=> s (triacylyglycerol production)
L1 0 (TRIACYLYGLYCEROL PRODUCTION)

=> s (triacylglycerol) and production
L2 1133 (TRIACYLGLYCEROL) AND PRODUCTION

=> s 12 and increase (oil content)
MISSING OPERATOR 'INCREASE (OIL'
The search profile that was entered contains terms or
nested terms that are not separated by a logical operator.

=> s 12 and (increase oil content)
L3 Q L2 AND (INCREASE OIL CONTENT)

=> s 12 and (enzyme)
I.4 175 I.2 AND (ENZYME)

=> s 14 and (catalyze an acyl-CoA independent reaction)
I-5 0 I-4 AND (CATALYZE AN ACYL-COA INDEPENDENT REACTION)

=> s 14 and (S. cerevisiae)

=> d 16 ti abs jbib tot

L6 ANSWER 1 OF 4 MEDLINE on STN
TI Synthesis of novel lipids in *Saccharomyces cerevisiae* by heterologous expression of an unspecific bacterial acyltransferase.
AB The bifunctional wax ester synthase/acyl-coenzyme A:diacylglycerol acyltransferase (WS/DGAT) is the key enzyme in storage lipid accumulation in the gram-negative bacterium *Acinetobacter calcoaceticus* ADP1, mediating wax ester, and to a lesser extent, triacylglycerol (TAG) biosynthesis. *Saccharomyces cerevisiae* accumulates TAGs and sterol esters as storage lipids. Four genes encoding a DGAT (*Dg1p*), a phospholipid:diacylglycerol acyltransferase (*Lro1p*) and two acyl-coenzyme

A:sterol acyltransferases (ASATs) (Are1p and Are2p) are involved in the final esterification steps in TAG and steryl ester biosynthesis in this yeast. In the quadruple mutant strain *S. cerevisiae* H1246, the disruption of DGA1, LRO1, ARE1, and ARE2 leads to an inability to synthesize storage lipids. Heterologous expression of WS/DGAT from *A. calcoaceticus* ADP1 in *S. cerevisiae* H1246 restored TAG but not steryl ester biosynthesis, although high levels of ASAT activity could be demonstrated for WS/DGAT expressed in *Escherichia coli* XL1-Blue in radiometric in vitro assays with cholesterol and ergosterol as substrates. In addition to TAG synthesis, heterologous expression of WS/DGAT in *S. cerevisiae* H1246 resulted also in the accumulation of fatty acid ethyl esters as well as fatty acid isoamyl esters. In vitro studies confirmed that WS/DGAT is capable of utilizing a broad range of alcohols as substrates comprising long-chain fatty alcohols like hexadecanol as well as short-chain alcohols like ethanol or isoamyl alcohol. This study demonstrated the highly unspecific acyltransferase activity of WS/DGAT from *A. calcoaceticus* ADP1, indicating the broad biocatalytic potential of this enzyme for biotechnological production of a large variety of lipids *in vivo* in prokaryotic as well as eukaryotic expression hosts.

ACCESSION NUMBER: 2004600773 MEDLINE
DOCUMENT NUMBER: PubMed ID: 15574908
TITLE: Synthesis of novel lipids in *Saccharomyces cerevisiae* by heterologous expression of an unspecific bacterial acyltransferase.
AUTHOR: Kalscheuer Rainer; Luftmann Heinrich; Steinbuchel Alexander
CORPORATE SOURCE: Institut fur Molekulare Mikrobiologie und Biotechnologie, Westfälische Wilhelms-Universität, Münster, Germany.
SOURCE: Applied and environmental microbiology, (2004 Dec) Vol. 70, No. 12, pp. 7119-25.
Journal code: 7605801. ISSN: 0099-2240.
PUB. COUNTRY: United States
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
LANGUAGE: English
FILE SEGMENT: Priority Journals
ENTRY MONTH: 200502
ENTRY DATE: Entered STN: 3 Dec 2004
Last Updated on STN: 16 Feb 2005
Entered Medline: 15 Feb 2005

L6 ANSWER 2 OF 4 MEDLINE on STN
TI Effect of hypertension on lipid metabolism and beta-oxidation in rat aorta and brain microvessels.
AB The effects of hypertension on various enzyme activities involved in lipid metabolism and beta-oxidation in rat brain microvessels and aorta were studied. The purity of the brain microvessel preparation was confirmed immunologically and microscopically. Activities involved in lipid synthesis, such as triacylglycerol synthesizing activity, acyl-CoA synthesizing activity, acyl-CoA: cholesterol acyltransferase and cytidine diphosphate choline:1,2-diacylglycerol cholinephosphotransferase, were significantly higher in brain microvessels than in aorta in both normotensive and hypertensive rats; lipid hydrolyzing activities, such as lipases and cholesterol esterases, were similar in the two preparations. Beta-oxidation in brain microvessels was more active than in aorta in both groups. Hypertension did not alter these enzyme activities in either aorta or brain microvessels, or change beta-oxidation in the aorta. However beta-oxidation in brain microvessels was significantly lower in hypertensive rats than in normotensive rats. These results suggest that brain microvessels are metabolically more active than aorta, and that their beta-oxidation activity is more susceptible to effects of hypertension. Reduced beta-oxidation in brain microvessels might lead to angioneurosis by derangement of energy production, which in turn may cause cerebral bleeding.

ACCESSION NUMBER: 83203441 MEDLINE

DOCUMENT NUMBER: PubMed ID: 6303275
TITLE: Effect of hypertension on lipid metabolism and beta-oxidation in rat aorta and brain microvessels.
AUTHOR: Sasaki N; Morisaki N; Shinomiya M; Matsuoka N; Saito Y;
Wakashin M; Ueda S; Kumagai A
SOURCE: Artery, (1982) Vol. 11, No. 2, pp. 108-18.
Journal code: 7508494. ISSN: 0098-6127.
PUB. COUNTRY: United States
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
LANGUAGE: English
FILE SEGMENT: Priority Journals
ENTRY MONTH: 198306
ENTRY DATE: Entered STN: 18 Mar 1990
Last Updated on STN: 6 Feb 1998
Entered Medline: 23 Jun 1983

L6 ANSWER 3 OF 4 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
TI Synthesis of novel lipids in *Saccharomyces cerevisiae* by heterologous expression of an unspecific bacterial acyltransferase.
AB The bifunctional wax ester synthase/acyl-coenzyme A:diacylglycerol acyltransferase (WS/DGAT) is the key enzyme in storage lipid accumulation in the gram-negative bacterium *Acinetobacter calcoaceticus* ADP1, mediating wax ester, and to a lesser extent, triacylglycerol (TAG) biosynthesis. *Saccharomyces cerevisiae* accumulates TAGs and steryl esters as storage lipids. Four genes encoding a DGAT (*Dga1p*), a phospholipid:diacylglycerol acyltransferase (*Lro1p*) and two acyl-coenzyme A:sterol acyltransferases (ASATs) (*Are1p* and *Are2p*) are involved in the final esterification steps in TAG and steryl ester biosynthesis in this yeast. In the quadruple mutant strain *S. cerevisiae* H1246, the disruption of *DGA1*, *LRO1*, *ARE1*, and *ARE2* leads to an inability to synthesize storage lipids. Heterologous expression of WS/DGAT from *A. calcoaceticus* ADP1 in *S. cerevisiae* H1246 restored TAG but not steryl ester biosynthesis, although high levels of ASAT activity could be demonstrated for WS/DGAT expressed in *Escherichia coli* XL1-Blue in radiometric in vitro assays with cholesterol and ergosterol as substrates. In addition to TAG synthesis, heterologous expression of WS/DGAT in *S. cerevisiae* H1246 resulted also in the accumulation of fatty acid ethyl esters as well as fatty acid isoamyl esters. In vitro studies confirmed that WS/DGAT is capable of utilizing a broad range of alcohols as substrates comprising long-chain fatty alcohols like hexadecanol as well as short-chain alcohols like ethanol or isoamyl alcohol. This study demonstrated the highly unspecific acyltransferase activity of WS/DGAT from *A. calcoaceticus* ADP1, indicating the broad biocatalytic potential of this enzyme for biotechnological production of a large variety of lipids in vivo in prokaryotic as well as eukaryotic expression hosts.

ACCESSION NUMBER: 2005:119076 BIOSIS
DOCUMENT NUMBER: PREV200500117462
TITLE: Synthesis of novel lipids in *Saccharomyces cerevisiae* by heterologous expression of an unspecific bacterial acyltransferase.
AUTHOR(S): Kalscheuer, Rainer; Luftmann, Heinrich; Steinbuechel, Alexander [Reprint Author]
CORPORATE SOURCE: Inst Mol Mikrobiol and Biotechnol, Univ Munster, Corrensstr 3, D-48149, Munster, Germany
steinbu@uni-muenster.de
SOURCE: Applied and Environmental Microbiology, (December 2004) Vol. 70, No. 12, pp. 7119-7125. print.
ISSN: 0099-2240 (ISSN print).
DOCUMENT TYPE: Article
LANGUAGE: English
ENTRY DATE: Entered STN: 23 Mar 2005
Last Updated on STN: 23 Mar 2005

L6 ANSWER 4 OF 4 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
TI Expression of Rhizopus oryzae lipase gene in Saccharomyces cerevisiae.
AB The extracellular production of active Rhizopus oryzae lipase (ROL) was carried out by the expression of the ProROL gene encoding a pro-form of ROL (ProROL) using prepro-alpha-factor in Saccharomyces cerevisiae. Two forms of recombinant ROL (rROL), rProROL by the expression of the ProROL gene and r28ROL which was a processed form of rProROL in the prosequence, were produced. Such a processing of rROL was catalyzed by the Kex2 membrane-bound endoprotease (Kex2p) in the late Golgi compartment. The ProROL and r28ROL could be produced independently as a single protein by the Kex2-engineered S. cerevisiae. Comparison of the properties of purified rROL showed that the prosequence modified some properties of ROL, and implied that the prosequence might play an physiologically important role in vivo. When only mature ROL (mROL) without the prosequence fused to the pre-alpha-factor leader sequence was expressed in S. cerevisiae, the enzyme activity was not observed in both the medium and cells. However, when the mROL was co-expressed in trans with the prosequence fused to the pre-alpha-factor leader sequence, the activity was recovered. The results showed that the prosequence may facilitate the folding of mROL, and the covalent linkage of the prosequence to the mROL was not necessary for the function. As the result of the deletion analysis at the N-terminus in the prosequence, the prosequence might work as an intramolecular chaperone. By the cell surface engineering using the gene encoding the C-terminal half of yeast alpha-agglutinin and the insertion of linker peptides, a novel strain displaying lipase on the cell surface was also constructed. Although S. cerevisiae itself is unable to utilize triolein, the transformant strain could grow on triolein as the sole carbon source. The cell surface-engineered yeast displaying ROL might be used as a potent biocatalyst.

ACCESSION NUMBER: 2002:401114 BIOSIS
DOCUMENT NUMBER: PREV200200401114
TITLE: Expression of Rhizopus oryzae lipase gene in Saccharomyces cerevisiae.
AUTHOR(S): Ueda, Mitsuyoshi [Reprint author]; Takahashi, Shouji; Washida, Motohisa; Shiraga, Seizaburo; Tanaka, Atsuo
CORPORATE SOURCE: Laboratory of Applied Biological Chemistry, Department of Synthetic Chemistry and Biological Chemistry, Graduate School of Engineering, Kyoto University, Sakyo-ku, Kyoto, 606-8501, Japan
miueda@sbchem.kyoto-u.ac.jp
SOURCE: Journal of Molecular Catalysis B Enzymatic, (7 June, 2002)
Vol. 17, No. 3-5, pp. 113-124. print.
ISSN: 1381-1177.
DOCUMENT TYPE: Article
General Review; (Literature Review)
LANGUAGE: English
ENTRY DATE: Entered STN: 24 Jul 2002
Last Updated on STN: 29 Aug 2002

=> d his

(FILE 'HOME' ENTERED AT 15:51:36 ON 15 JUN 2007)

FILE 'MEDLINE, BIOSIS' ENTERED AT 15:51:56 ON 15 JUN 2007

L1 0 S (TRIACYLYGLYCEROL PRODUCTION)
L2 1133 S (TRIACYLGLYCAROL) AND PRODUCTION
L3 0 S L2 AND (INCREASE OIL CONTENT)
L4 175 S L2 AND (ENZYME)
L5 0 S L4 AND (CATALYZE AN ACYL-COA INDEPENDENT REACTION)
L6 4 S L4 AND (S. CEREVISIAE)

=> s 12 and (DNA)
L7 52 L2 AND (DNA)

=> s 17 and 16
L8 0 L7 AND L6

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EPO Abstracts Database
JPO Abstracts Database
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IBM Technical Disclosure Bulletins

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DB=USPT; PLUR=YES; OP=OR

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|-----------|------------------------------|--------|-----------|
| <u>L3</u> | L2 and l1 | 5 | <u>L3</u> |
| <u>L2</u> | (triacylglycerol production) | 841118 | <u>L2</u> |
| <u>L1</u> | dahlqvist.in. | 23 | <u>L1</u> |

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Search Results - Record(s) 1 through 5 of 5 returned.

1. Document ID: US 6791008 B1

L3: Entry 1 of 5

File: USPT

Sep 14, 2004

US-PAT-NO: 6791008

DOCUMENT-IDENTIFIER: US 6791008 B1

TITLE: Use of a class of enzymes and their encoding genes to increase the oil content in transgenic organisms

DATE-ISSUED: September 14, 2004

INVENTOR-INFORMATION:

| NAME | CITY | STATE | ZIP CODE | COUNTRY |
|---------------------------|------------|-------|----------|---------|
| Banas; Antoni | Siedlce | | | PL |
| Sandager; Line | Copenhagen | | | DK |
| St.ang.hl; Ulf | Uppsala | | | SE |
| <u>Dahlqvist</u> ; Anders | Furulund | | | SE |
| Lenman; Marit | Lund | | | SE |
| Ronne; Hans | Uppsala | | | SE |
| Stymne; Sten | Svalov | | | SE |

US-CL-CURRENT: 800/281, 435/224, 435/471, 435/483, 536/23.1, 536/23.2, 536/23.7,
800/278, 800/298, 800/306

[Full](#) [Title](#) [Citation](#) [Front](#) [Review](#) [Classification](#) [Date](#) [Reference](#) [Sequences](#) [Attachments](#) [Claims](#) [KOMC](#) [Drawn D.](#)

2. Document ID: US 6524900 B2

L3: Entry 2 of 5

File: USPT

Feb 25, 2003

US-PAT-NO: 6524900

DOCUMENT-IDENTIFIER: US 6524900 B2

TITLE: Method concerning a junction barrier Schottky diode, such a diode and use thereof

DATE-ISSUED: February 25, 2003

INVENTOR-INFORMATION:

| NAME | CITY | STATE | ZIP CODE | COUNTRY |
|--------------------------|-------------|-------|----------|---------|
| <u>Dahlqvist</u> ; Fanny | Johanneshov | | | SE |

| | | |
|--------------------|--------------|----|
| Lendenmann; Heinz | Stocksund | SE |
| Hermannsson; Willy | Vaster.ang.s | SE |

US-CL-CURRENT: 438/167; 257/E21.359, 257/E27.051, 257/E29.104, 257/E29.338,
438/237, 438/328, 438/431

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Drawn D](#)

3. Document ID: US 6333448 B1

L3: Entry 3 of 5

File: USPT

Dec 25, 2001

US-PAT-NO: 6333448

DOCUMENT-IDENTIFIER: US 6333448 B1

TITLE: Plant enzyme and use thereof

DATE-ISSUED: December 25, 2001

INVENTOR-INFORMATION:

| NAME | CITY | STATE | ZIP CODE | COUNTRY |
|--------------------|--------------------|-------|----------|---------|
| Bafor; Maureen | Benin City | | | NG |
| Banas; Antoni | 08-110 Siedlce | | | PL |
| Dahlgvist; Anders | S-244 66 Furuland | | | SE |
| Gummeson; Per-Olov | S-227 38 Lund | | | SE |
| Lee; Michael | S-231 97 Klagstorp | | | SE |
| Sjodal; Staffan | S-756 50 Uppsala | | | SE |
| Stymne; Sten | S-268 90 Svalov | | | SE |
| Lenman; Marit | S-22359 Lund | | | SE |

US-CL-CURRENT: 800/295; 435/254.1, 435/255.1, 435/419, 435/69.1, 536/23.6, 800/281

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Drawn D](#)

4. Document ID: US 6306357 B1

L3: Entry 4 of 5

File: USPT

Oct 23, 2001

US-PAT-NO: 6306357

DOCUMENT-IDENTIFIER: US 6306357 B1

** See image for Certificate of Correction **

TITLE: Process and apparatus for absorbing hydrogen sulphide

DATE-ISSUED: October 23, 2001

INVENTOR-INFORMATION:

| NAME | CITY | STATE | ZIP CODE | COUNTRY |
|----------------|-------|-------|----------|---------|
| Simonson; Erik | Vaxjo | | | SE |

| | | |
|------------------------|--------------|----|
| Wallin; Mats | Lund | SE |
| Bengtsson; Sune | Vaxjo | SE |
| <u>Dahlqvist; Erik</u> | Vaster.ang.s | SE |

US-CL-CURRENT: 423/232; 162/51, 422/169, 422/170, 422/171, 422/181, 423/220

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Drawn D.](#)

5. Document ID: US PP08319 P

L3: Entry 5 of 5

File: USPT

Jul 27, 1993

US-PAT-NO: PP08319

DOCUMENT-IDENTIFIER: US PP08319 P

TITLE: Poinsettia plant `Lilo White`

DATE-ISSUED: July 27, 1993

INVENTOR-INFORMATION:

| | | | | |
|--------------------------------|-----------|-------|----------|---------|
| NAME | CITY | STATE | ZIP CODE | COUNTRY |
| <u>Dahlqvist; Kjell-Ingvar</u> | Hollviken | | | SE |

US-CL-CURRENT: PLT/304

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Drawn D.](#)

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